

(FILE 'HOME' ENTERED AT 16:34:28 ON 10 APR 2000)

FILE 'MEDLINE, BIOSIS, EMBASE, CAPLUS' ENTERED AT 16:34:40 ON 10 APR 2000

L1 895 S ADENINE NUCLEOTIDE TRANSLOCATOR

L2 184 S ANT1

L3 10 S ANT1 AND FUSION

L4 4 DUP REM L3 (6 DUPLICATES REMOVED)

L5 2 S ANT1 AND FUSION

L6 106 S ANT2

L7 59 DUP REM L6 (9 DUPLICATES REMOVED)

L8 7 S L7 AND MEMBRANE

L9 97 S ANT2 AND EXPRESS?

L10 22 S ANT2 AND CDNA

L11 8 DUP REM L10 (14 DUPLICATES REMOVED)

L12 7 S ANT2 AND PURIFICATION

L13 113 S ANT 3

L14 9 S ANT3 AND PURIFICATION

L15 9 S ANT3 AND PURIF?

L16 4 S ANT3 AND CDNA

L17 9 S ANT3 5A EXPRESS?

L18 9 S ANT3 5A PROTEIN

L19 21 S L1 AND FUSION

L20 9 DUP REM L19 (12 DUPLICATES REMOVED)

EAST

	TYP	L	Hit	Text	Search	DBS	Time	Stamp
e	#	s						
1	ERS	L2	1	and defining a 1 nucleotide adj transcript	UCFAT; EIG; PERWENT; 10/04/10 17:11			
2	ERS	L2	1	and defining a 1 nucleotide adj transcript	UCFAT; EIG; PERWENT; 10/04/10 17:11			
3	ERS	L2	1	and defining a 1 nucleotide adj transcript	UCFAT; EIG; PERWENT; 10/04/10 17:11			
3	ERS	L2	3	and defining a 1 nucleotide adj transcript	UCFAT; EIG; PERWENT; 10/04/10 17:11			
4	ERS	L4	2	and defining a 1 nucleotide adj transcript	UCFAT; EIG; PERWENT; 10/04/10 17:11			

LB ANSWER 7 OF 7 CAPLUS COPYRIGHT 2000 ACS
AN 1397:673204 CAPLUS
DN 127:342467
TI Somatic cell mapping of the adenine nucleotide translocator gene family
IN cattle
AU Li, Lei; Womack, James E.
CS Department Veterinary Pathobiology, Texas A&M University, College
Station,
TX, 77843, USA
SO Mamm. Genome (1997), 8(10), 773-774
CODEN: MAMGEC; ISSN: 0938-8990
PB Springer
PT Journal
LA English
AB Adenine nucleotide translocator [ADP/ATP translocase, (ANT), or ADP/ATP
carrier (AAC)] is the most abundant mitochondrial protein. As an
integral
component of the inner mitochondrial **membrane**, it catalyzes the
exchange of intramitochondrial ATP for cytoplasmic ADP, consequently
controlling the ATP supply of the cell. Its central role in cellular
energy supply suggests that ANT might be regulated in different tissues
TC
for tissue specific functional and developmental requirements. The
authors assigned the bovine adenine nucleotide translocator ANT3 gene to
chromosome X which segregated concordantly with **ANT2**. Both
ANT2 and ANT3 have been mapped on human chromosome X. However,
ANT3 escapes X inactivation on the pseudautosomal region of Xp22.3 in
human, whereas **ANT2** is subjected to X-inactivation and localized
on Xq13-q26 (Chen et al. 1990). Further localization of these genes will
help clarify the evolutionary history of mammalian sex chromosomes.

LA ANSWER 6 OF 7 CAPLUS COPYRIGHT 2000 ACS
AN 1999:696536 CAPLUS
TI Stress sensitive B encodes an adenine nucleotide translocase in
Drosophila
 melanogaster
AU Zhang, Yong Q.; Root, John; Brogna, Saverio; Davis, Andrew W.; Barbash, Daniel A.; Nash, David; Ashburner, Michael
J Department of Genetics, University of Cambridge, Cambridge, CB2 3EH, UK
S Genetics (1999), 153(2), 391-393
CODEN: GENTAE; ISSN: 0016-6731
PB Genetics Society of America
PT Journal
LA English
AF Adenine nucleotide translocases (ANT) are required for the exchange of
ADP
 and ATP across the inner mitochondrial **membrane**. They are
 essential for life, and most eukaryotes have at least two different Ant
 genes. Only one gene had been described from *Drosophila*, and this had
 not
 been characterized genetically. We show that mutations in this gene
 correspond to the previously described loci, *sesB* and *l(l)9Ed*.
 Immediately adjacent to this gene is another encoding a second ANT
 protein, which has 78% identity to that encoded by *sesB/l(l)9Ed*. These
 two genes are transcribed from a common promoter, and their mRNAs are
 produced by differential splicing. Hutter and Karch suggested that the
 sesB ANT gene corresponded to *Hmr*, a gene identified by an allele that
 rescues otherwise inviable interspecific hybrids between *Drosophila*
 melanogaster and its sibling species. This hypothesis is not supported
 by

1.8 ANSWER 4 OF 7 MEDLINE
AN 92340491 MEDLINE
DN 92340491
TI Differential expression of adenine nucleotide translocator isoforms in mammalian tissues and during muscle cell differentiation.
AU Stepien G; Torrcini A; Chung A B; Hodge J A; Wallace D C
DS Department of Genetics and Molecular Medicine, Emory University School of Medicine, Atlanta, Georgia 30322..
NC HL-45572 (NHLBI)
NS-21328 (NINDS)
JO JOURNAL OF BIOLOGICAL CHEMISTRY, (1992 Jul 25) 267 (21) 14592-7.
Journal code: HIV. ISSN: 0021-9258.
CY United States
PT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals; Cancer Journals
EM 199210
AB The adenine nucleotide translocator (ANT) catalyzes the exchange of ADP and ATP across the mitochondrial internal **membrane**. Its three isoforms, **ANT1**, **ANT2**, and **ANT3** are coded by differentially regulated nuclear genes. The patterns of expression of these genes in human, bovine, and mouse tissue are similar. **ANT1** is highly expressed in heart and skeletal muscle and is induced during myoblast differentiation. It is coordinately regulated with the nuclear gene for the mitochondrial ATP synthase beta subunit, with which it shares the positive muscle cis element, the **CXBOX**. **ANT2** is either absent or weakly expressed in all tissues. **ANT3** is ubiquitously expressed in all tissues, and its transcript level is proportional to the level of oxidative metabolism.
The tissue-specific expression of the ANT gene family thus provides insight into the molecular basis of the differential reliance of mammalian tissues on oxidative phosphorylation.

L11 ANSWER 7 OF 8 BIOSIS COPYRIGHT 2000 BIOSIS
AN 1994:435865 BIOSIS
DN PREV199497448865
TI A human pseudoautosomal gene ADP'ATP translocase, escapes X-inactivation whereas a homologue on Xq is subject to X-inactivation.
AU Schiebel, Katrin (1); Weiss, Birgit (1); Woehrle, Doris; Rappold, Gudrun (1)
CS (1) Institut Human Genetics, Univ. of Heidelberg, D-6900 Heidelberg Germany
SO Nature Genetics, (1993) Vol. 3, No. 2, pp. 81-87.
ISSN: 1061-4036.
PT Article
LA English
AB We report the cloning of a highly conserved pseudoautosomal gene on the human sex chromosomes. A **cDNA** clone was selected by crosshybridization with a microdissected clone from the chromosomal subregion Xp22.3. It encodes a previously characterized member of the ADP'ATP translocase family and plays a fundamental role in cellular energy metabolism. This gene, **ANT3**, is located approximately 1,300 kilobases from the telomere, proximal to the pseudoautosomal gene **CSPRA**, and escapes X-inactivation. Interestingly, a homologue of **ANT3**, **ANT2**, maps to Xq and is subject to X-inactivation. These genes provide the first evidence of two closely related X-chromosomal genes, which show striking differences in their X-inactivation behaviour.

L20 ANSWER 2 OF 9 MEDLINE
AN 97284663 MEDLINE
DN 97284663
TI Thyroid hormone activates transcription from the promoter regions of some human nuclear-encoded genes of the oxidative phosphorylation system.
AU Li R; Liciakova K; Zaid A; Betina S; Frideil E; Nelson B D
CS Department of Biochemistry, Stockholm University, Sweden.
SO MOLECULAR AND CELLULAR ENDOCRINOLOGY, (1997 Apr 4) 128 (1-2): 69-75.
Journal code: E69. ISSN: 0303-7207.
CY Ireland
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 199710
EW 19971002
AB Thyroid hormone (T3) modulates the mRNA levels for cytochrome c1 and the **adenine nucleotide translocator-2 (ANT2)** in adult rat liver. Here we show that T3 activates expression of a reporter gene driven from the human cytochrome c1 and ANT2 promoters transfected into human choriocarcinoma JEG3 cells. By contrast, the human F1-ATPase beta-subunit promoter responded marginally, thus providing a pattern of differential expression similar to that earlier observed in rats *in vivo*. T3-activation is dependent on co-expression of the thyroid hormone receptor (TR alpha1). Co-expression of both the TR and RXR receptors had no additional effect. Transient transfection of deletion constructs showed that T3 activation is retained by the proximal regions of the cytochrome c1 and ANT2 promoters, and, in the case of cytochrome c1, is lost upon removal of a fragment containing the transcription initiator ((nucleotides) (nt) + 1 tc + 100). The promoter regions supporting T3-activation of the reporter genes appear to lack strong DNA binding